

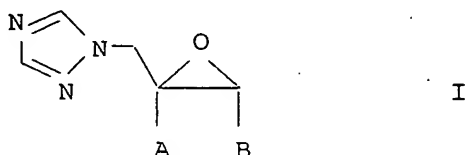
Preparation 1,2,4-triazolylmethyloxiranes

Abstract

5

The present invention relates to a process for the preparation of 1,2,4-triazol-1-ylmethyloxiranes of the formula I

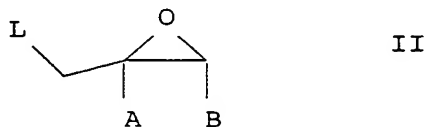
10



15 in which A and B are identical or different and, independently of one another, are C<sub>1</sub>-C<sub>4</sub>-alkyl, phenyl-C<sub>1</sub>-C<sub>2</sub>-alkyl, C<sub>3</sub>-C<sub>6</sub>-cycloalkyl, C<sub>3</sub>-C<sub>6</sub>-cycloalkenyl, tetrahydropyranyl, tetrahydrofuranyl, dioxanyl or phenyl, where the phenyl radical can carry one to three substituents chosen from the group: halogen, nitro, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-alkyloxy, phenoxy, amino, C<sub>1</sub>-C<sub>2</sub>-haloalkyl or phenylsulfonyl, which comprises reacting

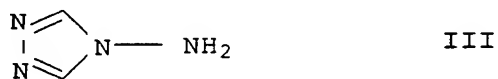
a) an oxirane of the formula II

25

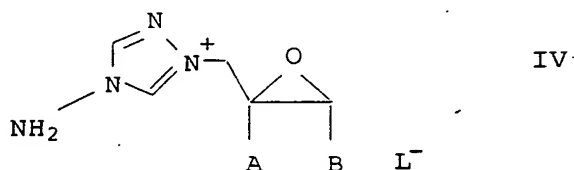


30 in which A and B have the meanings given above and L is a nucleophilically substitutable leaving group, with 4-amino-1,2,4-triazole of the formula III

35



40



45 to give 4-amino-1,2,4-triazolium salts of the formula IV and

20

b.) deaminating the 4-amino-1,2,4-triazolium salts IV with alkali metal nitrites and acid or organic nitrites to give 1,2,4-triazol-1-ylmethyloxiranes of the formula I,

5 and to 4-aminotriazolium salts of the formula IV as intermediates.

10

15

20

25

30

35

40

45